(Item 1 from file: 350) DIALOG(R) File 350: Derwent WPIX (c) 2003 Thomson Derwent. All rts. reserv. 014659303 **Image available** WPI Acc No: 2002-480007/200251 XRPX Acc No: N02-379059 Removable fixing device for neuro implants, especially retina implants, has fixing head that allows implant to be removed from beneath it Patent Assignee: TD VERW GMBH (TDTD-N); INTELLIGENT IMPLANTS GMBH (INTE-N) Inventor: ECKMILLER R Number of Countries: 089 Number of Patents: 003 Patent Family: Patent No Kind Date: Applicat No Kind Date Week 20020606 WO 200243631 A2 WO 2001EP14077 Α 20011203 200251 DE 10060029 Α1 20020613 DE 1060029 Α 20001201 200251 AU 200219150 20020611 AU 200219150 Α Α 20011203 200264 Priority Applications (No Type Date): DE 1060029 A 20001201 Patent Details: Patent No Kind Lan Pg Filing Notes Main IPC WO 200243631 A2 G 13 A61F-009/00 Designated States (National): AE AL AM AT AU AZ BA BB BG BR BY CA CH CN CU CZ DK EE ES FI GB GD GE GH GM HU ID IL IN IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MA MD MG MK MN MW MX NO NZ PL PT RO RU SD SE SG SI SK SL TJ TM TR TT TZ UA UG US UZ VN YU ZA ZW Designated States (Regional): AT BE CH CY DE DK EA ES FI FR GB GH GM GR IE IT KE LS LU MC MW MZ NL OA PT SD SE SL SZ TR TZ UG ZM ZW DE 10060029 A61F-002/14 Α1 AU 200219150 A A61F-009/00 Based on patent WO 200243631 Abstract (Basic): WO 200243631 A2 NOVELTY - The head of the fixing device (3) extends over the implant (2) on the opposite side to the retina. The fixing device can be removed during the re-explanation or fixing process by pulling the microcontact foil beneath the head. DETAILED DESCRIPTION - INDEPENDENT CLAIMS are also included for (a) a fixing device with a head which extends over the implant surface and which can be pivoted, folded or pulled out in order to release the implant, and (b) a fixing device with a head which can be releasably secured in place by an anchor structure extending through the retina, pigment epithel and vein wall. USE - None given. ADVANTAGE - Neuro implants can be releasably fixed to retina tissue, enabling a defective implant to be replaced, or a more up to date implant to be inserted in place of the old one. DESCRIPTION OF DRAWING(S) - Figure 1 shows a section of tissue from the retina with a microcontact film lying epiretinally on top of it, held in place by pivotable fixing devices. Retina tissue section (1) Microcontact film (2) Fixing device (3) Pivot arm (4) Release position of pivot arm (5) Movement of pivot arm into release position (6) pp; 13 DwgNo 1/3 Title Terms: REMOVE; FIX; DEVICE; NEURO ; IMPLANT; RETINA; IMPLANT; FIX; HEAD; ALLOW; IMPLANT; REMOVE; BENEATH Derwent Class: P32

International Patent Class (Main): A61F-002/14; A61F-009/00

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(Item 2 from file: 350)
DIALOG(R) File 350: Derwent WPIX
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014172377
             **Image available**
WPI Acc No: 2001-656605/200175
XRPX Acc No: N01-489484
 Micro-contact structure for neuro -prostheses for implantation on nerve
  tissue has multiple contacts on two dimensional carrier panel which can
Patent Assignee: INTELLIGENT IMPLANTS GMBH (INTE-N); TD VERW GMBH (TDTD-N);
  BECKER M (BECK-I); ECKMILLER R (ECKM-I); HUNERMANN R (HUNE-I)
Inventor: BECKER M ; ECKMILLER R ; HUENERMANN R; HUNERMANN R
Number of Countries: 031 Number of Patents: 006
Patent Family:
Patent No
              Kind
                     Date
                              Applicat No
                                             Kind
                                                    Date
                                                              Week
                    20011101 US 2001771283
US 20010037061 A1
                                              Α
                                                   20010126
                                                              200175
DE 10020846
               Α1
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                             DE 1020846
                                                  20000428
                                                             200203
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WO 200183025
               Α1
                   20011108
                             WO 2000EP12713
                                                  20001214
                                              Α
                                                             200212
AU 200131589
                   20011112
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                                                  20001214
               Α
                                              Α
                                                             200222
EP 1276537
               A1 20030122
                             EP 2000991189
                                              Α
                                                  20001214
                                                             200308
                              WO 2000EP12713 A
                                                  20001214
TW 478933
                   20020311
                             TW 2000127777
                                              Α
                                                  20010201
                                                             200309
Priority Applications (No Type Date): DE 1020846 A 20000428
Patent Details:
Patent No Kind Lan Pg
                         Main IPC
                                      Filing Notes
US 20010037061 A1
                      8 A61B-005/04
DE 10020846
                       A61F-002/02
              Α1
WO 200183025 A1 G
                       A61N-001/05
   Designated States (National): AU BR CA IL JP KR MX NZ SG US
   Designated States (Regional): AT BE CH CY DE DK ES FI FR GB GR IE IT LU
   MC NL PT SE TR
AU 200131589 A
                       A61N-001/05
                                      Based on patent WO 200183025
EP 1276537
           . A1 G
                       A61N-001/05
                                      Based on patent WO 200183025
   Designated States (Regional): AT BE CH CY DE DK ES FI FR GB GR IE IT LI
   LU MC NL PT SE TR
TW 478933
                       A61F-002/00
              Α
Abstract (Basic): US 20010037061 A1
        {\tt NOVELTY - The \; micro-contact \; structure \; for \quad } \textbf{neuro -} \textbf{prostheses} \; \; \textbf{has}
    multiple contacts formed on a two dimensional carrier which has at
    least two regions which can move relative to one another. The regions
    can assume a base position and an operating position. The size of the
    micro-contact structure is collapsed during surgical transportation to
    the implant point my moving the relatively movable sections.
        USE - For implantation at mammalian muscle tissue, or blood vessels
    or body organs
        ADVANTAGE - Allows ease of positioning implant
        DESCRIPTION OF DRAWING(S) - Drawing shows plan view of implant
        pp; 8 DwgNo 1/4
Title Terms: MICRO; CONTACT; STRUCTURE; NEURO; PROSTHESIS; IMPLANT; NERVE
  ; TISSUE; MULTIPLE; CONTACT; TWO; DIMENSION; CARRY; PANEL; CAN; FOLD
Derwent Class: P31; P32; P34
International Patent Class (Main): A61B-005/04; A61F-002/00; A61F-002/02;
  A61N-001/05
International Patent Class (Additional): A61F-002/14
File Segment: EngPI
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(Item 3 from file: 350) DIALOG(R) File 350: Derwent WPIX (c) 2003 Thomson Derwent. All rts. reserv. 014141156 **Image available** WPI Acc No: 2001-625367/200172 XRPX Acc No: N01-466125 Secure operating method for neuro -prosthesis in central nervous system within scull, by performing data transmission when authorisation signal transmitted from external to internal components is checked and accepted Patent Assignee: INTELLIGENT IMPLANTS GMBH (INTE-N); TD VERW GMBH (TDTD-N) Inventor: BECKER M ; ECKMILLER R ; HUENERMANN R; ORTMANN V Number of Countries: 030 Number of Patents: 005 Patent Family: Date Applicat No Kind Patent No Kind Date WO 200147598 Α1 20010705 WO 2000EP6666 Α 20000713 200172 B 20010709 AU 200068231 20000713 AU 200068231 Α Α 200172 20010906 DE 19962915 A1 DE 1062915 Α 19991223 200172 US 6493587 В1 20021210 US 2000635016 20000809 Α 200301 EP 1307258 A1 20030507 EP 2000956178 20000713 200332 Α WO 2000EP6666 20000713 Α Priority Applications (No Type Date): DE 1062915 A 19991223 Patent Details: Main IPC . Patent .No Kind Lan Pg Filing Notes WO 200147598 A1 G 42 A61N-001/36 Designated States (National): AU BR CA CN IL JP KR MX NZ SG US Designated States (Regional): AT BE CH CY DE DK ES FI FR GB GR IE IT LU MC NL PT SE AU 200068231 A61N-001/36 Based on patent WO 200147598 DE 19962915 A61F-002/00 Α1 US 6493587 A61N-001/37 В1 EP 1307258 Al G A61N-001/36 Based on patent WO 200147598 Designated States (Regional): AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU MC NL PT SE Abstract (Basic): WO 200147598 A1 NOVELTY - At least one neuro -prosthesis component is implanted so that it is in contact with a nerve tissue or is associated with a nerve tissue in such a way that they interact. The neuro -prosthesis is only operated during the period of specific authorisation, and/or the system comprises an authorised data transmission between external components and implanted components, and/or an authorised communication for monitoring and/or fixing the neuro -prosthesis operating status, and/or the communication between the external and implanted components is encrypted. USE - None given. ADVANTAGE - Prevents unauthorised access to data. DESCRIPTION OF DRAWING(S) - The drawing shows a protection system for a neuro -prosthesis.

SYSTEM; SCULL; PERFORMANCE; DATA; TRANSMISSION; AUTHORISE; SIGNAL; TRANSMIT; EXTERNAL; INTERNAL; COMPONENT; CHECK; ACCEPT Derwent Class: P34; S05; T01; W02 International Patent Class (Main): A61F-002/00; A61N-001/36; A61N-001/37 International Patent Class (Additional): A61N-001/372 File Segment: EPI; EngPI

Title Terms: SECURE; OPERATE; METHOD; NEURO; PROSTHESIS; CENTRAL; NERVE;

pp; 42 DwgNo 1/5

(Item 4 from file: 350) DIALOG(R) File 350: Derwent WPIX (c) 2003 Thomson Derwent. All rts. reserv. 013967307 **Image available** WPI Acc No: 2001-451521/200148 XRAM Acc No: C01-136298 Treatment of physiological disorders requiring anticoagulation therapy by inhibition of Factor Xa with a sulfonic acid or sulfonylamino N-(heteroaralkyl)-azaheterocyclylamide derivative Patent Assignee: AVENTIS PHARMA DEUT GMBH (AVET) Inventor: BARTON J N; BECKER M R; CHOI-SLEDESKI Y M; EWING W R; GONG Y; GREEN D M; LEVELL J; PAULS H W Number of Countries: 093 Number of Patents: 002 Patent Family: Patent No Kind Date Applicat No Kind Date Week WO 200139759 A2 20010607 WO 2000EP11577 A 20001121 200148 B AU 200117042 20010612 AU 200117042 Α 20001121 Priority Applications (No Type Date): US 99453307 A 19991202 Patent Details: Patent No Kind Lan Pg Main IPC Filing Notes WO 200139759 A2 E 106 A61K-031/00 Designated States (National): AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CR CU CZ DE DK DM DZ EE ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MA MD MG MK MN MW MX MZ NO NZ PL PT RO RU SD SE SG SI SK SL TJ TM TR TT TZ UA UG UZ VN YU ZA ZW Designated States (Regional): AT BE CH CY DE DK EA ES FI FR GB GH GM GR IE IT KE LS LU MC MW MZ NL OA PT SD SE SL SZ TR TZ UG ZW AU 200117042 A A61K-031/00 Based on patent WO 200139759 Abstract (Basic): WO 200139759 A2 NOVELTY - A method for treating physiological disorders by inhibition of Factor Xa comprises administration of a diagnostic, cardioprotective, direct thrombin inhibiting, anticoagulant, antiplatelet or fibrinolytic agent with a sulfonic acid or sulfonylamino N-(heteroaralkyl)-azaheterocyclylamide derivative (I). DETAILED DESCRIPTION - A method for treating physiological disorders by inhibition of Factor Xa comprises administration of a diagnostic, cardioprotective, direct thrombin inhibiting, anticoagulant, antiplatelet or fibrinolytic agent with a sulfonic acid or sulfonylamino N-(heteroaralkyl)-azaheterocyclylamide derivative of formula (I) or its salt or N-oxide. Ar1=heteroaryl; Z'=alkylenyl or substituted amide or amine derivative; R1, R2, X1, X1a, X3, X4=H or optionally substituted (hetero) hydrocarbyl; X2, X2a=H or=0; and X5, X5a, X5b=amino, amido or sulfonamide derivative. INDEPENDENT CLAIMS are also included for: (1) compositions comprising a diagnostic, cardioprotective, direct thrombin inhibiting, anticoagulant, antiplatelet or fibrinolytic agent and a compound of formula (I); and.

MECHANISM OF ACTION - Factor Xa inhibition. 7-methoxynaphthalene-2-sulfonic acid

ACTIVITY - Anticoagulant; cardiant; antianginal; thrombolytic; vasotropic; cerebroprotective; immunosuppressive; virucide; cytostatic;

(2) a kit for carrying out the method.

neuroprotective

(1-(1,6-diaminoisoquinolin-7-ylmethyl)-2-oxopyrrolidin-3-yl)-amide trifluoroacetate (Ia) showed a Ki of 80 nM.

USE - The method is useful for treating physiological disorders by inhibition of Factor Xa, including abnormal thrombus formation, acute myocardial infarction, unstable angina, thromboembolism, acute vessel closure associated with thrombolytic therapy or percutaneous transluminal coronary angioplasty, transient ischemic attacks, stroke, pathologic thrombus formation occurring in the veins of the lower extremities following abdominal, knee or hip surgery, risk of pulmonary thromboembolism or disseminated systemic intravascular coagulopathy occurring in vascular systems during septic shock, certain viral infections or cancer.

pp; 106 DwgNo 0/0

Title Terms: TREAT; PHYSIOLOGICAL; DISORDER; REQUIRE; ANTICOAGULANT;

THERAPEUTIC; INHIBIT; FACTOR; ACID; N; DERIVATIVE

Derwent Class: B02; B03; D16

International Patent Class (Main): A61K-031/00

File Segment: CPI

10/5/5 (Item 5 from file: 350)

DIALOG(R) File 350: Derwent WPIX

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013707132 **Image available**
WPI Acc No: 2001-191356/200119

Related WPI Acc No: 2000-086312; 2000-422942

XRAM Acc No: C01-057286

New oxoazaheterocyclyl compounds useful are factor Xa inhibitors used for treating thrombotic complications and chronic and degenerative diseases
Patent Assignee: AVENTIS PHARM PROD INC (AVET); AVENTIS PHARM INC (AVET)

Inventor: **BECKER M R**; BURNS C J; CHOI-SLEDESKI Y M; CONDON S M; DAVIS R S; 'EWING W R; HANNEY B A; HE W; JIANG J Z; LAU W F; LI A; MYERS M R; PAULS H W; POLI G B; SPADA A P

Number of Countries: 094 Number of Patents: 008

Patent Family:

racene ramary.									
Pat	tent No	Kind	Date	App	plicat No	Kind	Date	Week	
WO	200107436	A2	20010201	WO	2000IB1156	A	20000726	200119	В
ΑU	200064628	Á	20010213	ΑU	200064628	A	20000726	200128	
BR	200013179	Α	20020402	BR	200013179	A	20000726	200231	
		,		WO	2000IB1156	Α	20000726		
CZ	200200323	А3	20020515	WO	2000IB1156	A	20000726	200241	
				CZ	2002323	Α	20000726		
ΕP	1208097	A2	20020529	EP	2000951781	Α	20000726	200243	
				WO	2000IB1156	Α	20000726		
SK	200200118	A3	20021106	WO	2000IB1156	A	20000726	200281	
				SK	2002118	A	20000726		
HU	200203375	A2	20021228	WO	2000IB1156	A	20000726	200308	
				HU	20023375	Α	20000726		
JΡ	2003508353	W	20030304	WO	2000IB1156	A	20000726	200319	
				JΡ	2001512520	Α	20000726		

Priority Applications (No Type Date): US 99363196 A 19990728 Patent Details:

Patent No Kind Lan Pg Main IPC Filing Notes WO 200107436 A2 E 459 C07D-403/06

Designated States (National): AE AG AL AM AT AU AZ BA BB BG BR BY CA CH CN CR CU CZ DE DK DM DZ EE ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MA MD MG MK MN MW MX MZ NO NZ PL PT RO RU SD SE SG SI SK SL TJ TM TR TT TZ UA UG US UZ VN YU ZA ZW

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Designated States (Regional): AT BE CH CY DE DK EA ES FI FR GB GH GM GR
   IE IT KE LS LU MC MW MZ NL OA PT SD SE SL SZ TZ UG ZW
                       C07D-403/06
AU 200064628 A
                                     Based on patent WO 200107436
                                     Based on patent WO 200107436
BR 200013179 A
                       C07D-403/06
CZ 200200323 A3.
                       C07D-403/06
                                     Based on patent WO 200107436
EP 1208097
             A2 E
                       C07D-403/06
                                     Based on patent WO 200107436
   Designated States (Regional): AL AT BE CH CY DE'DK ES FI FR GB GR IE IT
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                       C07D-403/06
                                     Based on patent WO 200107436
SK 200200118 A3
HU 200203375 A2
                                     Based on patent WO 200107436
                       C07D-403/06
JP 2003508353 W
                   722 C07D-241/08
                                     Based on patent WO 200107436
Abstract (Basic): WO 200107436 A2
        NOVELTY - Oxoazaheterocyclyl compounds (I) are new.
        DETAILED DESCRIPTION - Oxoazaheterocyclyl compounds of formula (I)
    and their salts, prodrugs, N-oxides, hydrates and solvates are new.
        G1, G2=L1-Cy1 or L2-Cy2;
        Cy1, Cy2=aryl, heteroaryl, cycloalkyl, cycloalkenyl, heterocyclyl,
    heterocyclenyl, fused arylcycloalkyl, fused aryl cycloalkenyl, fused
    aryl heterocyclyl, fused aryl heterocyclenyl, fused
    heteroarylcycloalkyl, fused heteroarylcycloalkenyl, fused heteroaryl
    heterocyclyl or fused heteroaryl heterocyclenyl (all optionally
    substituted);
        L1=absent, O, NR5, S(P)p, S(O)pNR5, CXY, L3-Q-L4-Q'-L5, COYCXY,
    CXYCO, CONR5S(O)p or COCONR5S(O)p;
        L2=absent or (CR7R8)q-Z-(CR9R10);
        L3, L5=absent or alkylene, alkenylene or alkynylene (all optionally
    substituted);
        L4=alkylene, alkenylene or alkynylene (all optionally substituted);
        Q, Q'=absent, O, S, NR5, S(O)p, S(O)pNR5 or CXY;
        A=CH or N;
        R1, R1a, R2, R2a, R3, R3a, R4, R4a=H, carboxy, alkoxycarbonyl,
    YlY2NCO, or alkyl, aryl, aralkyl, heteroaryl or heteroaralkyl (all
    optionally substituted), or
        R1 + R1a, R2 + R2a, R3 + R3a or R4 + R4a=0 or S, or
        CR1R2=cycloalkyl, cycloalkenyl, heteroaryl or heteroaralkyl (all
    optionally substituted) or cycloalkyl, cycloalkenyl, heterocyclyl, or
    heterocyclenyl; or
        CR3R4=cycloalkyl, cycloalkenyl, heterocyclyl or heterocyclenyl, or
        R1a, R2a=absent and
        CR1R2=aryl or heteroaryl, or
        R3a, R4a=absent and
        CR3R4=aryl or heteroaryl or
        one or more of the pairs CR1R1a, CR2R2a, CR3R3a or CR4R4a=3-7
    membered cycloalkyl or cycloalkenyl, or
        m, n=0-2, provided that m and n are not both 0 and provided that
    when R1 + R1a=0 or S, n=1 and when R4 + R4a=0 or S, m=1;
        R5=H, R6O(CH2)v, R6O2C(CH2)x, Y1Y2NCO(CH2)x, Y1Y2N(CH2)v or alkyl,
    aralkyl or heteroaralkyl (all optionally substituted);
        R6=H or alkyl, aralkyl or heteroaralkyl (all optionally
    substituted);
        Y1, Y2=H or alkyl, alkoxy, aryloxy, aryl, aralkyl or heteroaralkyl
    (all optionally substituted); or
        NY1Y2=monocyclic heterocyclyl;
        R7-R10=H, OH, alkoxy or alkyl, aryl, heteroaryl, aralkyl or
    heteroaralkyl (all optionally substituted); provided that only one of
    R7 and R8 or R9 and R10=alkoxy or hydroxy and provided that when any of
    R7-R10=hydroxy or alkoxy, then the hydroxy or alkoxy is not
    alpha-substituted to an N, O or S in Z;
        X=0 or S;
        Y=absent, O, S or NR5;
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Z=absent, optionally substituted lower alkenylene or alkynylene or O, CO, S(O)p, NR5, NR5CO or CONR5;

x=1-4;

v=2-4;

p=1 or 2 and

q, r=0-3, provided that both are not 0;

L1=O, NR5, S(O)p, S(O)pNR5, CXY or L3-Q-L4-Q'-L5;

provided that:

- (1) L1=and R3 + R3a=O or S, then R2, R2a=H, alkyl, aminoalkyl, alkylaminoalkyl, alkoxy, alkoxyalkyl, alkoxyaminoalkyl, cycloalkyl alkylamino, benzyloxyalkyl, isopropyl, amino methyl, methoxyethyl aminomethyl, piperazine, pyrrolidine, ethoxymethyl, benzyloxymethyl, methoxymethyl, isobutyl, isopropylamino or isopropylaminomethyl; provided that R2 and R2a are not both H;
- (2) when L1=0, NR5, S(O)p, S(O)pNR5, CXY or L3-Q-L4-Q'-L5 and R3 + R3a=O or S, then R4 + R4a=O or S;
- (3) when L1=O, NR5, S(O)p, S(O)pNR5, CXY or L3-Q-L4-Q'-L5 and R3 + R3a=O or S, then Cy1=thiophen-isoxazole, thiophene-pyrazole, thiophene-oxadiazole, thiophene-thiadiazole, thiophene-triazole, thiophene-pyridine or phenyl-triazole and Cy2=aminoquinazoline or pyrrolo-pyridine;
- (4) when L1=O, NR5, S(O)p, S(O)pNR5, CXY or L3-Q-L4-Q'-L5, then CR1R2=cycloalkyl, cycloalkenyl, heterocyclyl or heterocyclenyl; or CR3R4=cycloalkyl, cycloalkenyl, heterocyclyl or heterocyclenyl; or R1a and R2a are absent and CR1R2=aryl or heteroaryl; or R3a and R4a are absent and CR3R4=aryl or heteroaryl; or one or more of the pairs CR1R1a, CR2R2a, CR3R3a or CR4R4a=3-7 membered cycloalkyl or cycloalkenyl;
- (5) when L1=0, NR5, S(0)p, S(0)pNR5, CXY or L3-Q-L4-Q'-L5, then R1, R1a, R2, R2a, R3, R3a, R4, R4a=Y1Y2NCO and Y1, Y2=H, or optionally substituted alkoxy or aryloxy, but Y1 and Y2 are not both H or
- (6) when L1=O, NR5, S(O)p, S(O)pNR5, CXY or L3-Q-L4-Q'-L5, then Z=CO and
- (7) when R1 and R1a or R4 and R4a=0 or S, then G1=L2-Cy2 and G2=L1-Cy1, or
- (8) when $\dot{R}2$ and $\dot{R}2a$ or $\dot{R}3$ and $\dot{R}3a=0$ or S, then G1=L1-Cy1 and G2=L2-Cy2.

ACTIVITY - Anticoagulant; thrombolytic; antirheumatic; antiarthritic; cytostatic; antiarteriosclerotic; nootropic. neuroprotective.

Activity tests, such as experimental in vivo rabbit venous thrombosis model, are described but no results are given.

MECHANISM OF ACTION - Factor Xa inhibitor.

USE - Useful for treating thrombotic complications. Inhibition of factor Xa is useful not only in anticoagulation therapy but whenever inhibition of blood coagulation is required such as to prevent coagulation of stored blood. Factor Xa inhibitors may also be used for the treatment or prevention of chronic and degenerative diseases such as arthritis, cancer, atherosclerosis and Alzheimer's disease.

pp; 459 DwgNo 0/0

Title Terms: NEW; COMPOUND; USEFUL; FACTOR; INHIBIT; TREAT; THROMBUS; COMPLICATED; CHRONIC; DEGENERATE; DISEASE

Derwent Class: B02; B03

International Patent Class (Main): C07D-241/08; C07D-403/06
International Patent Class (Additional): A61K-031/495; A61K-031/496;
A61K-031/498; A61K-031/502; A61K-031/506; A61K-031/517; A61K-031/519;
A61K-031/5377; A61P-007/02; A61P-009/02; A61P-009/04; A61P-009/10;
A61P-019/02; A61P-025/28; A61P-035/00; A61P-041/00; A61P-043/00;
C07D-401/06; C07D-401/14; C07D-403/14; C07D-405/14; C07D-409/06;
C07D-409/12; C07D-409/14; C07D-413/14; C07D-417/14; C07D-471/04;
C07D-491/04; C07D-495/04

NO 200003808 A

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10/5/6
            (Item 6 from file: 350)
DIALOG(R) File 350: Derwent WPIX
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             **Image available**
WPI Acc No: 2000-086312/200007
Related WPI Acc No: 2000-422942; 2001-191356
XRAM Acc No: C00-023997
  New heterocyclyl compounds are factor Xa inhibitors useful for treating
  unstable angina, stroke, etc.
Patent Assignee: AVENTIS PHARM PROD INC (AVET ); RHONE-POULENC RORER PHARM
  INC (RHON ); AVENTIS PHARM INC (AVET )
Inventor: BECKER M R ; BURNS C J; CHOI-SLEDESKI Y M; CONDON S M; DAVIS R S
  ; EWING W R; HANNEY B A; HE W; JIANG J Z; LAU W F; LI A; MYERS M R; PAULS
  H W; POLI G B; SPADA A P; JIANG J; CHOISLEDESKI Y M; DAVIES R S
Number of Countries: 082 Number of Patents: 014
Patent Family:
              Kind
                      Date
                              Applicat No
                                              Kind
                                                     Date
Patent No
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WO 9937304
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                              WO 99US1682
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SK 200001108
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JP 2002501024
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                              JP 2000528286
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AU 745425
                    20020321
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                                                   19990127
                                                              200233
HU 200101810
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                                                   19990127
                                                              200238
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                              HU 20011810
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                              WO 99US1682
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MX 2000007250
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Priority Applications (No Type Date): US 9872707 P 19980127
Patent Details:
Patent No Kind Lan Pg
                          Main IPC
                                       Filing Notes
              A1 E 298 A61K-031/505
WO 9937304
   Designated States (National): AL AM AT AU AZ BA BB BG BR BY CA CN CU CZ
   DE DK EE ES FI GB GE GH HU IL IS JP KE KG KP KR KZ LC LK LR LS LT LU LV
   MD MG MK MN MW MX NO NZ PL PT RO RU SD SE SG SI SK SL TJ TM TR TT UA UG
   US UZ VN YU ZW
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                    349 A61K-000/00
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BR 9907300
                        A61K-031/505
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                                      Based on patent WO 9937304
EP 1051176
                        A61K-031/505 Based on patent WO 9937304
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   Designated States (Regional): AL AT BE CH DE DK ES FI FR GB GR IE IT LI
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C07C-409/12

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                       A61K-031/505
              B
                                     Previous Publ. patent AU 9926533
                                     Based on patent WO 9937304
HU 200101810 A2
                       C07C-409/14
                                     Based on patent WO 9937304
NZ 505960
              Α
                       C07D-409/14
                                     Based on patent WO 9937304
MX 2000007250 A1
                       A61K-031/505
Abstract (Basic): WO 9937304 A1
        NOVELTY - Novel substituted oxoazaheterocyclyl compounds are factor
    Xa inhibitors useful for treating acute myocardial infarction, thrombus
    formation and cancer
        DETAILED DESCRIPTION - Substituted oxoazaheterocyclyl compounds of
    formula (I) and their salts, prodrugs, N-oxides, hydrates and solvates
        G1,G2=L1-Cy1 or L2-Cy2; provided that when R1 and R1a or R4 and
    R4a=0 or S, then G1=L2-Cy2 and G2=L1-Cy1, or when R2 + R2a R3 and R3a=0
    or S, then G1=L1-Cy, and G2=L2-Cy2;
        Cy1, Cy2=aryl, heteroaryl, cycloalkyl, cycloalkenyl, heterocyclyl,
    heterocyclenyl, fused aryl cycloalkyl, fused arylcycloalkenyl, fused
    arylheterocyclyl, fused aryl heterocyclenyl, fused
    heteroarylcycloalkyl, fused heteroaryl cycloalkenyl, fused
    heteroarylheterocyclyl or fused heteroaryl heterocyclenyl (all
    optionally substituted);
        L1=O, NR5, S(O)p, S(O)pNR5, C(X)Y or L3-Q-L4-Q'-L2;
        L3, L5=a bond or alkylene, alkenylene or alkynylene (all optionally
    substituted;
        L4=alkylene, alkenylene or alkynylene (all optionally substituted);
        Q, Q'=absent, O, S, NR5, S, S(O)pNR5 or C(X)Y;
        A=CH or N;
        R1-R4, R1a-R4a=H, carboxy, alkoxycarbonyl, Y1Y2NCO or alkyl, aryl,
    aralkyl, heteroaryl or heteroaralkyl (all optionally substituted); or
        R1 + R1a, R2 + R2a, R3 + R3a, R4 + R4a=0 or S;
        m, n=0-2; provided that both are not O; that when R1 + R1a=O or S,
    n=1; and when R4 + R4a=O or S, m=1;
        L2=absent or (CR7R8)q-Z-(CR9R10)r;
        R5=H, R6O(CH2)v, R6O2C(CH2)x, Y1Y2NCO(CH2)x, Y1Y2N(CH2)v or alkyl,
    aralkyl or heteroaralkyl (all optionally substituted);
        R6=H or alkyl, aralkyl, or heteroaralkyl (all optionally
    substituted);
        Y1, Y2=H or alkyl, aryl, aralkyl or heteroaralkyl (all optionally
    substituted); or
        NY1Y2=monocyclic heterocyclyl;
        R7-R10=H, OH, alkoxy or alkyl, aryl, heteroaryl, aralkyl or
    heteroaralkyl (all optionally substituted); provided that only one of
    R7 and R8 or one of R9 and R10=OH, or alkoxy; and that when R7-R10=or
    alkoxy, then the OH or alkoxy is not alpha-substituted to N, O or S in
    Z;
        X=0 or S;
        Y=absent, O, S or NR5;
        Z=absent, O, S(O)p, NR5, NR5CO, CONR5 or lower alkenylene or lower
    alkynylene (both optionally substituted);
        x=1-4;
        v=2-4;
        p=1 or 2;
        q, r=0-3; provided that both are not 0.
        An INDEPENDENT CLAIM is included for intermediates of formula (II).
        R1'-R4', R1a'-R4a'=H, carboxy, alkoxycarbonyl, Y1Y2NCO or alkyl,
    aryl, aralkyl, heteroaryl or heteroaralkyl (all optionally
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substituted); P=H or N-protecting group L2' = (CR7R8) q - Z - (CR9R10) r. ACTIVITY - Vasotropic; Thrombolytic; Anticoagulant; Antianginal; Cardiant; Cerebroprotective; Immunosuppressive; Antibacterial; Virucide; Cytostatic; Antiarthritic; Antiarteriosclerotic; Neuroprotective ; Nootropic. MECHANISM OF ACTION - Factor-X-Inhibitor (claimed). Both the activity of free Factor Xa and Factor Xa assembled in the prothrombinase complex (Factor, Xa, Factor Va, calcium and phospholipid) and inhibited by (I). Assays are described, but no results are given. USE - (I) are useful for treating conditions capable of being modulated by inhibiting activity of Factor Xa such as venous vasculature, arterial vasculature, abnormal thrombus formation, acute myocardial infarction, unstable angina, thromboembolism, acute vessel closure associated with thrombolytic therapy, percutaneous transluminal coronary angioplasty, transient ischemic attacks, stroke, intermittent claudication or bypass grafting of the coronary or peripheral arteries, vessel luminal narrowing, restenosis post coronary or venous angioplasty, maintenance of vascular access patency in long-term hemodialysis patients, pathologic thrombus formation occurring in the veins of the lower extremities following abdominal, knee and hip surgery, a risk of pulmonary thromboembolism, or disseminated systemic intravascular coagulopathy occurring in vascular systems during septic shock, certain viral infections or cancer (claimed). (I) may also be used for treating chronic and degenerative diseases such as arthritis, atherosclerosis and Alzheimer's disease and abnormal proliferation of cells. pp; 298 DwgNo 0/0 Title Terms: NEW; HETEROCYCLE; COMPOUND; FACTOR; INHIBIT; USEFUL; TREAT; UNSTABLE; ANGINA; STROKE Derwent Class: B02; B03 International Patent Class (Main): A61K-000/00; A61K-031/505; C07C-409/12; C07C-409/14; C07D-401/12; C07D-409/14; C07D-419/00 International Patent Class (Additional): A61K-031/4365; A61K-031/437; A61K-031/495; A61K-031/496; A61K-031/517; A61P-007/00; A61P-007/02; A61P-009/10; A61P-019/02; A61P-025/28; A61P-035/00; A61P-043/00; C07C-401/12; C07D-241/18; C07D-401/04; C07D-401/14; C07D-403/04; C07D-403/06; C07D-403/14; C07D-405/04; C07D-409/04; C07D-409/06; C07D-409/12; C07D-413/14; C07D-417/14; C07D-441/08; C07D-471/04; C07D-491/04; C07D-495/04; C07D-519/00; C07D-471-04; C07D-495-04 File Segment: CPI (Item 7 from file: 350) DIALOG(R) File 350: Derwent WPIX (c) 2003 Thomson Derwent. All rts. reserv. 012851474 WPI Acc No: 2000-023306/200002 XRAM Acc No: C00-005678 Improved preparation of carbocyclic compounds Patent Assignee: GILEAD SCI INC (GILE-N); SOUTHCO INC (SOUT-N) Inventor: BECKER M W ; CHAPMAN H H; KELLY D E; KENT K M; LEW W; LOUIE M S; MCGEE L R; POSTICH M J; PRISBE E J; ROHLOFF J C; SCHULTZE L M; YU R H; ZHANG L Number of Countries: 086 Number of Patents: 003

Patent Family: Patent No

WO 9955664

Kind

A1

Date

Applicat No

19991104 WO 99US7378

Kind

Α

Date

Week

19990423 200002 B

19991116 AU 9936376 AU 9936376 19990423 200015 Α Α 20010425 KR 2000713006 KR 2001034875 A 20001120 200164 Α Priority Applications (No Type Date): US 9882994 P 19980424; US 9882994 A 19980522; US 99274858 A 19990323 Patent Details: Patent No Kind Lan Pq Main IPC Filing Notes A1 E 69 C07C-247/14 WO 9955664 Designated States (National): AE AL AM AT AU AZ BA BB BG BR BY CA CH CN CU CZ DE DK EE ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MD MG MK MN MW MX NO NZ PL PT RO RU SD SE SG SI SK SL TJ TM TR TT UA UG US UZ VN YU ZA ZW Designated States (Regional): AT BE CH CY DE DK EA ES FI FR GB GH GM GR IE IT KE LS LU MC MW NL OA PT SD SE SL SZ UG ZW AU 9936376 Based on patent WO 9955664 KR 2001034875 A F16B-021/00 Abstract (Basic): WO 9955664 Al NOVELTY - Process comprises one, or sequential combinations, of preparations of compounds. DETAILED DESCRIPTION - Process comprises preparations (14) of 3,4-isopropylidenel,5-quinic lactone (305 or 100); (15) ethyl-(-)-3,4-isopropylidene quinate (307 or 101); (16) ethyl-(-)-3,4-isopropylidene-5-O-methanesulfonyl quinate (308 or 102); (17) ethyl-3,4-0-isopropylidene-5-0-methanesulfonyl Shikimate (309 or

DETAILED DESCRIPTION - Process comprises preparations (14) of 3,4-isopropylidene1,5-quinic lactone (305 or 100); (15) ethyl-(-)-3,4-isopropylidene quinate (307 or 101); (16) ethyl-(-)-3,4-isopropylidene-5-O-methanesulfonyl quinate (308 or 102); (17) ethyl-3,4-O-isopropylidene-5-O-methanesulfonyl Shikimate (309 or 104); (18) ethyl-3,4-O-isopropylidene-5-O-methanesulfonyl Shikimate (300 or 107); (19) ethyl-3-O-(1-ethylpropyl)-5-O-methansulfonyl Shikimate (301 or 108); (20) ethyl (3R,4R,5S)-4,5-epoxy-3-(1-ethylpropoxy)-1-cyclohexane-1-carboxylate (304 or 110); (21) ethyl (3R,4R,5S)-5-azido-3-(1-ethylpropoxy)-4-hydroxy-cyclohexene-1-carboxylate (313 or 112); (22) ethyl (3R,4R,5S)-4,5-imino-3-(1-ethylpropoxy)-1-cyclohexane-1-carboxylate (315 or 113); (23) ethyl (3R,4R,5S)-4-amino-5-azido-3-(1-ethylpropoxy)-1-cyclohexene-1-carboxyla

te (316 or 114); (24) ethyl (3R,4R,5S)-4-acetamido-5-azido-3-(1-ethylpropoxy)-1-cyclohexene-1-carbo xylate (317 or 115) and/or (25) ethyl (3R,4R,5S)-4-acetamido-5-azido-3-(1-ethylpropoxy)-1-cyclohexene-1-carbo xylate phosphate (1:1) (116.H3PO4).

USE - Used for preparing carbocyclic compounds. Used to prepare intermediates for synthesis of neuraminidase inhibitors. Used to prepare products useful as monomers for synthesis of polymers with unique pendent functionalities, including polyamides, polyesters and mixed polyester-polyamides, for use as cation exchange agents, in preparation of molecular sieves, textiles, fibers, films and formed articles, as well as to prepare polyfunctional surfactants with useful surfactant, surface coating, emulsion modifying, rheology modifying and surface-wetting properties. Used to produce polyfunctional compounds with defined geometries carrying simultaneously polar and non-polar groups for use as phase-transfer agents for phase-transfer catalysis and liquid/liquid ion extraction. Used to produce compounds optionally containing asymmetric carbon atoms for use as chiral auxiliaries in synthesis or resolution of optically active materials such as in resolution of carboxylic acid racemic mixture. Used to produce linkers or spacers in preparing affinity absorption matrices, immobilized enzymes for process control or immunoassay reagents. Used to produce products for crosslinking with affinity reagents such as hormones, peptides, antibodies, drugs and insoluble substrates to absorb binding partners for affinity reagents from manufactured preparations,

diagnostic samples and other impure mixtures and to easily recover immobilized enzymes following catalytic conversions.

ADVANTAGE - Improves preparations of (100), (101), (102), (104), (107), (108), (110), (111), (113), (115), (116), phosphate salt of (116) and hydrochloride salt of (116) (claimed).

pp; 69 DwgNo 0/0

Title Terms: IMPROVE; PREPARATION; CARBOCYCLIC; COMPOUND

Derwent Class: B02; B05; Q47; Q61

International Patent Class (Main): C07C-247/14; F16B-021/00

International Patent Class (Additional): CO7D-317/48

File Segment: CPI; EngPI

10/5/8 (Item 8 from file: 350)

DIALOG(R) File 350: Derwent WPIX

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012126555 **Image available** WPI Acc No: 1998-543467/199847

XRPX Acc No: N98-423049

Apparatus for promoting selective stimulation of defective retina - employs portable laser and video prosthetic in form of conventional spectacles which cause laser beam to stimulate relevant neuron groups

Patent Assignee: BECKER M (BECK-I)

Inventor: BECKER M

Number of Countries: 001 Number of Patents: 001

Patent Family:

Patent No Kind Date Applicat No Kind Date Week
DE 19713612 A1 19981015 DE 1013612 A 19970402 199847 B

Priority Applications (No Type Date): DE 1013612 A 19970402

Patent Details:

Patent No Kind Lan Pg Main IPC Filing Notes

DE 19713612 A1 8 A61F-009/08

Abstract (Basic): DE 19713612 A

An apparatus for selectively stimulating one or a number of nerve cells in the defective retina of a human eye employs a portable laser source whose beam is projected onto the retina via the natural lens of the eye.

The laser unit and system controller can be conveniently carried in a small container at the waist with connections made by coaxial and fibre-optics cables to a prosthetic assembly in the form of conventional spectacles.

These incorporate a photosensor array for detecting the wearer's forward environment, a fixed focusing mirror and a beam steering reflector responding to signals from a pupil position sensor which combine to regulate stimulation of the appropriate retinal neurons .

USE - Provides retinal stimulation in situations where disease has impaired efficient functioning.

ADVANTAGE - Is able to be more precise in terms of effect on individual cells or small groups of cells then current systems employing electrical stimulation via electrodes. Does not require expense and risks of invasive surgery which attends implantation of micro-photodiodes.

Dwg.2/4

Title Terms: APPARATUS; PROMOTE; SELECT; STIMULATING; DEFECT; RETINA; EMPLOY; PORTABLE; LASER; VIDEO; PROSTHESIS; FORM; CONVENTION; SPECTACLE; CAUSE; LASER; BEAM; STIMULATING; RELEVANT; NEURON; GROUP

Derwent Class: P32; P34; S05

International Patent Class (Main): A61F-009/08

International Patent Class (Additional): A61N-005/06

File Segment: EPI; EngPI

10/5/9 (Item 9 from file: 350)

DIALOG(R) File 350: Derwent WPIX

(c) 2003 Thomson Derwent. All rts. reserv.

012040911 **Image available** WPI Acc No: 1998-457821/199840

XRPX Acc No: N98-357358

Encoder for visual neuroprosthetic providing active vision. - uses adjustable receptive field characteristic filters inserted in signal path between photosensor array and implanted stimulation and registration interface

Patent Assignee: INTELLIGENT IMPLANTS GMBH (INTE-N); TD VERW GMBH (TDTD-N);

ECKMILLER R (ECKM-I); IIP TECHNOLOGIES GMBH (IIPT-N)

Inventor: ECKMILLER R

Number of Countries: 075 Number of Patents: 024

Patent Family:

Pat	ent No	Kind	Date	App	olicat No	Kind	Date	Week	
DE	19707046	A1	19980827	DE	1007046	A	19970221	199840	В
WO	9836793	A2	19980827	WO	98EP971	A	19980220	199840	
WO	9836795	A1	19980827	WO	98EP968	Α	19980220	199840	
WO	9837691	A1	19980827	WO	98EP970	Α	19980220	199840	
ΑU	9864990	Α	19980909	AU	9864990	Α	19980220	199905	
AU	9867223	Α	19980909	AU	9867223	A	19980220	199905	
ΑU	9868220	A	19980909	AU	9868220	Α	19980220	199905	
ΕP	969896	A2	20000112	ΕP	98913568	Α	19980220	200008	
				WO	98EP971	A	19980220		
DE	19880174	. T	20000105	DE	1080174	A	19980220	200009	
				WO	98EP970	Α	19980220		
ΕP	971770	A1	20000119	EP	98912345	A	19980220	200009	
				WO	98EP968	A	19980220		
BR	9807260	Α	20000502	BR	987260	Α	19980220	200033	
				WO	98EP971	Α	19980220		
BR	9807847	Α	20000829	BR	987847	A	19980220	200046	
				WO	98EP968	Α	19980220		
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MX	9907732	A1	20000401	MX	997732	Α	19990820	200124	
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JP	2001523989	W	20011127		98536259	A	19980220	200204	
					98EP968	A	19980220		
US	6400989	В1	20020604	MO		Α	19980220	200242	
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NZ	337392	A	20020628	NZ	337392	A	19980220	200252	
	6500054				98EP971	Α	19980220		
US	6530954	В1	20030311	WO		A	19980220	200321	
				US	2000367138	A	20000229		

Priority Applications (No Type Date): DE 1007046 A 19970221 Patent Details:

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Patent No Kind Lan Pg
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WO 9836793
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WO 9836795
              A1 G
                       A61N-001/36
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   MN MW MX NO NZ PL PT RO RU SD SE SG SI SK SL TJ TT UA US UZ VN YU ZW
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WO 9837691
              A1 G
                       H04N-005/232
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AU 9864990
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                                     Based on patent WO 9837691
AU 9867223
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                       A61N-001/36
                                     Based on patent WO 9836795
AU 9868220
              Α
                       A61N-001/00
                                     Based on patent WO 9836793
EP 969896
              A2 G
                                     Based on patent WO 9836793
                       A61N-001/00
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DE 19880174
                       H04N-005/232
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              A1 G
                       A61N-001/36
                                     Based on patent WO 9836795
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                                     Based on patent WO 9836793
BR 9807847
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                       A61F-002/02
                                     Based on patent WO 9836795
MX 9907727
              A1
                       A61N-001/00
MX 9907732
              Α1
                       A61N-001/36
AU 732190
                       A61N-001/36
                                     Previous Publ. patent AU 9867223
                                     Based on patent WO 9836795
KR 2000075557 A
                       A61N-001/36
                                     Based on patent WO 9836795
                                     Based on patent WO 9836793
KR 2000075560 A
                       A61N-001/00
JP 2001511687 W
                    25 A61N-001/36
                                     Based on patent WO 9836793
JP 2001523989 W
                    34 A61F-009/08
                                     Based on patent WO 9836795
US 6400989
              В1
                       A61N-001/18
                                     Based on patent WO 9836795
AU 747686
              В
                       A61N-001/00
                                     Previous Publ. patent AU 9868220
                                     Based on patent WO 9836793
NZ 337366
              Α
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                                     Based on patent WO 9836795
                       A61N-001/36
NZ 337392
              Α
                                     Based on patent WO 9836793
US 6530954
              В1
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                                     Based on patent WO 9836793
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Abstract (Basic): DE 19707046 A

The encoder has the signals provided by a photosensor array fed to a stimulation and registration interface, provided by an implanted microcontact structure, via adjustable receptive field characteristic filters. The latter receive signals provided by the stimulation and registration interface, for providing an active vision function. The photosensor array may be incorporated in a spectacles frame, with image tracking movement controlled via head and eye movement detectors.

 $\ensuremath{\mathsf{USE}}$ - For $\ensuremath{\mathsf{neuroprosthetic}}$ retina implant for blind patient, for night vision etc.

ADVANTAGE - Autonomous object detection and following. Dwg.1/3

Title Terms: ENCODE; VISUAL; ACTIVE; VISION; ADJUST; RECEPTIVE; FIELD; CHARACTERISTIC; FILTER; INSERT; SIGNAL; PATH; PHOTOSENSOR; ARRAY; IMPLANT

```
; STIMULATING; REGISTER; INTERFACE
Derwent Class: P32; P34; S05; T01; W03; W04
International Patent Class (Main): A61F-002/02; A61F-009/08; A61N-001/00;
  A61N-001/18; A61N-001/36; G06F-017/00; H04N-005/232
International Patent Class (Additional): A61F-002/14; A61F-002/18;
  A61F-002/48; A61F-011/04; A61N-001/08; G05B-015/02; G06F-015/18;
  G06F-019/00; H04N-005/14
File Segment: EPI; EngPI
             (Item 10 from file: 349)
DIALOG(R) File 349: PCT FULLTEXT
(c) 2003 WIPO/Univentio. All rts. reserv.
00954399
            **Image available**
METHOD FOR WEAVING AN AIRBAG
PROCEDE DE TISSAGE D'UN SAC GONFLABLE
VERFAHREN ZUM WEBEN EINES LUFTSACKS
Patent Applicant/Assignee:
  BERGER SEIBA-TECHNOTEX VERWALTUNGS GMBH & CO, Ballyweg 5, 79713 Bad
    Sackingen, DE, DE (Residence), DE (Nationality), (For all designated
    states except: US)
Patent Applicant/Inventor:
  ESCHBACH Thomas, Rheinpromenade 10, 79790 Rheinheim, DE, DE (Residence),
    DE (Nationality), (Designated only for: US)
   BECKER Michael , Adelsberg 11, 79669 Zell im Wiesental, DE, DE
    (Residence), DE (Nationality), (Designated only for: US
Legal Representative:
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English Abstract
  Disclosed is a method for weaving a single-pieced airbag (2) or air tube
  consisting of at least two layers on a mechanical loom, characterized in
  that weft yarns of different strengths are woven in at least one layer.
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French Abstract

L'invention concerne un procede de tissage d'un sac gonflable (2) ou d'une chambre a air bicouche d'une seule piece sur un metier a tissier, ledit procede etant caracterise en ce que, dans une couche au moins, des fils de trame de grosseurs differentes sont tisses.

German Abstract

Es wird Verfahren zum Weben eines wenigstens zweilagigen, einstuckigen Luftsacks (2) oder Luftschlauchs auf einer Webmaschine vorgeschlagen, das durch gekennzeichnet ist, dass in wenigstens einer Lage Schussfaden unterschiedlicher Starken verwebt werden.

Legal Status (Type, Date, Text)
Publication 20021107 Al With international search report.
Publication 20021107 Al Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.

Examination 20021219 Request for preliminary examination prior to end of 19th month from priority date

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